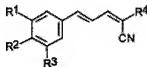


I. Amendments to the Claims

This listing of claims will replace all prior versions and listings of claims in the application.

Listing of Claims:

1. (Currently Amended) A compound of Formula I, or a salt, solvate, or hydrate thereof:



I

wherein

R¹ and R² are each independently selected from H, OH, C₁₋₆alkyl, C₁₋₆alkoxy, C₁₋₆alkylCO₂, NH₂, NH-C₁₋₆alkyl, N(C₁₋₆alkyl)(C₁₋₆alkyl), C₁₋₆alkyl(C=O)NH, C₁₋₆alkyl(C=O)N(C₁₋₆alkyl), SH, S-C₁₋₆alkyl, O-Si(C₁₋₆alkyl)(C₁₋₆alkyl)(C₁₋₆alkyl), NO₂, CF₃, OCF₃ and halo, or R¹ and R² together represent O-C₁₋₆alkyl-O, thereby forming a ring both OH or R¹ and R² are both OCH₃;

R³ is selected from H, OH, C₁₋₆alkyl, C₁₋₆alkoxy, C₁₋₆alkylCO₂, NH₂, NH-C₁₋₆alkyl, N(C₁₋₆alkyl)(C₁₋₆alkyl), C₁₋₆alkyl(C=O)NH, or C₁₋₆alkyl(C=O)N(C₁₋₆alkyl), SH, S-C₁₋₆alkyl, O-Si(C₁₋₆alkyl)(C₁₋₆alkyl)(C₁₋₆alkyl), NO₂, halo and CH₂-(CH₂)_n-Ar;

R⁴ is selected from C(X)R⁵, SO₃Ar, NH₂, NH-C₁₋₆alkyl, N(C₁₋₆alkyl)(C₁₋₆alkyl), P(O)(OH)₂, P(O)(OC₁₋₆alkyl)₂, and C(NH₂)=C(CN)₂;

X is selected from O, S, NH and N-C₁₋₆alkyl;

R⁵ is selected from NH₂, OH, NH(CH₂)_pAr, NH(CH₂)_pOH, (CH₂)_pOC₁₋₆alkyl, C₁₋₆alkyl, C₁₋₆alkoxy, NHNH₂, NHC(O)NH₂, NHC(O)C₁₋₆alkoxy, N-morpholino and N-pyrrolidino; and

Ar is an aromatic or heteroaromatic group, unsubstituted or substituted with 1-4 substituents, independently selected from OH, C₁₋₆alkyl, C₁₋₆alkoxy, NH₂, NH-C₁₋₆alkyl, N(C₁₋₆alkyl)(C₁₋₆alkyl), SH, S-C₁₋₆alkyl, NO₂, CF₃, OCF₃ and halo;

n is 0 to 4; and

p is 1-4 ;

provided that at least one of R¹, R², and R³ is selected from C₁₋₆alkylCO₂, C₁₋₆alkyl(C=O)NH, or C₁₋₆alkyl(C=O)N(C₁₋₆alkyl) or

~~R¹ and R² together represent O-C₁₋₆alkyl-O, thereby forming a ring.~~

2-5. (Cancelled)

6. (Currently Amended) The compound according to claim 1, wherein R³ is selected from H, OH, C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₄alkylCO₂, NH₂, NH-C₁₋₄alkyl, N(C₁₋₄alkyl)(C₁₋₄alkyl), C₁₋₄alkyl(C=O)NH, or C₁₋₄alkyl(C=O)N(C₁₋₄alkyl), ~~SH, S-C₁₋₄alkyl, NO₂ and halo.~~

7. (Currently Amended) The compound according to claim 6, wherein R³ is selected from H, OH, OCH₃, CH₃CO₂, SH, SMe, NO₂, CH₃CONH, or CH₃CONCH₃, ~~and halo.~~

8. (Cancelled)

9. (Previously Presented) The compound according to claim 1, wherein R⁴ is selected from C(X)R⁵ and C(NH₂)=C(CN)₂.

10. (Original) The compound according to claim 9, wherein R⁴ is C(X)R⁵.

11. (Previously Presented) The compound according to claim 10, wherein X is selected from O and S.

12. (Previously Presented) The compound according to claim 10, wherein R⁵ is selected from NH₂, OH, NH(CH₂)_pAr, NH(CH₂)_pOH and C₁₋₄alkoxy.

13. (Original) The compound according to claim 12, wherein p is 1-3.

14. (Currently Amended) The compound according to claim ~~13~~ 12, wherein R⁵ is selected from NH₂, OH, NH(CH₂)_pAr, NH(CH₂)_pOH and OCH₃.

15. (Original) The compound according to claim 14, wherein p is 1-2.

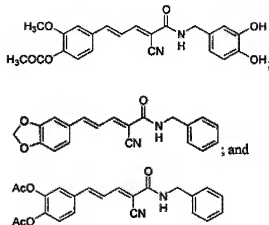
16. (Currently Amended) The compound according to claim 1, wherein Ar is an unsubstituted phenyl group or a phenyl group substituted with 1-4 substituents ~~optionally~~ independently selected from OH, C₁₋₆alkyl, C₁₋₆alkoxy, NH₂, NH-C₁₋₆alkyl, N(C₁₋₆alkyl)(C₁₋₆alkyl), SH, S-C₁₋₆alkyl, NO₂, CF₃, OCF₃ and halo.

17. (Currently Amended) The compound according to claim 14, wherein Ar is an unsubstituted phenyl group or a phenyl group substituted with 1-4 substituents ~~optionally~~ independently selected from OH, C₁₋₆alkyl, C₁₋₆alkoxy, NH₂, NH-C₁₋₆alkyl, N(C₁₋₆alkyl)(C₁₋₆alkyl), SH, S-C₁₋₆alkyl, NO₂, CF₃, OCF₃ and halo.

18. (Currently Amended) The compound according to any of claims 16 and 17, wherein Ar is an unsubstituted phenyl group or phenyl group substituted with 1-2 substituents ~~optionally~~ independently selected from OH, C₁₋₄alkyl, C₁₋₄alkoxy, NH₂, NH-C₁₋₄alkyl, N(C₁₋₄alkyl)(C₁₋₄alkyl), SH, S-C₁₋₄alkyl, NO₂, CF₃, OCF₃ and halo.

19. (Currently Amended) The compound according to claim 18, wherein Ar is an unsubstituted phenyl group or phenyl group substituted with 1-2 substituents ~~optionally~~ independently selected from OH, OCH₃, NH₂, NHCH₃, N(CH₃)₂, SH, SCH₃, CF₃, OCF₃ and halo.

20. (Previously Presented) A compound selected from:



21. (Currently Amended) A composition comprising a compound according to claim 1 or claim 38 in admixture with a pharmaceutically acceptable diluent or carrier.

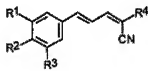
22-29 (Cancelled)

30. (Currently Amended - Withdrawn) A method of modulating cell proliferation comprising administering an effective amount of a compound capable of modulating cell proliferation according to claim 1 or claim 38 or a composition according to claim 21 to a cell or animal in need thereof.

31. (Currently Amended - Withdrawn) A method of inhibiting cell proliferation comprising administering an effective amount of a compound capable of inhibiting cell proliferation according to claim 1 or claim 38 or a composition according to claim 21 to a cell or animal in need thereof.

32. (Currently Amended - Withdrawn) A method of inhibiting cancer cell proliferation comprising administering an effective amount of a compound capable of inhibiting cancer cell proliferation according to claim 1 or claim 38 or a composition according to claim 21 to a cell or animal in need thereof.

33. (Currently Amended - Withdrawn) A method of treating cancer comprising administering an effective amount of a compound capable of inhibiting cancer cell proliferation according to claim 1 or claim 38 or a composition according to claim 21 to a cell or animal in need thereof.
34. (Currently Amended - Withdrawn) A method according to claim 32 or 33 wherein said cancer is a hematopoietic cell cancer.
35. (Currently Amended - Withdrawn) A method according to claim 32 or 33 wherein said cancer is a leukemia, a lymphoma, a myeloma or a carcinoma.
36. (Previously Presented - Withdrawn) A method according to claim 35 wherein said leukemia is acute lymphoblastic leukemia, aggressive Philadelphia+ leukemia, acute myelocytic leukemia, chronic myeloid leukemia, chronic lymphocytic leukemia or juvenile myelomonocyte leukemia,
37. (Previously Presented - Withdrawn) A method according to claim 35 wherein said leukemia is acute lymphoblastic leukemia.
38. (New) A compound of Formula I, or a salt, solvate, or hydrate thereof:



wherein

R^1 is OCH_3 ; and R^2 is OH;

R^3 is selected from $C_{1-6}alkylCO_2$, $C_{1-6}alkyl(C=O)NH$, or $C_{1-6}alkyl(C=O)N(C_{1-6}alkyl)$;

R^4 is selected from $C(X)R^5$, SO_3Ar , NH_2 , $NH-C_{1-6}alkyl$, $N(C_{1-6}alkyl)(C_{1-6}alkyl)$, $P(O)(OH)_2$, $P(O)(OC_{1-6}alkyl)_2$, and $C(NH_2)=C(CN)_2$;

X is selected from O, S, NH and N- $C_{1-6}alkyl$;

R^5 is selected from NH_2 , OH, $NH(CH_2)_pAr$, $NH(CH_2)_pOH$, $(CH_2)_pOC_{1-6}alkyl$, $C_{1-6}alkyl$, $C_{1-6}alkoxy$, $NHNH_2$, $NHC(O)NH_2$, $NHC(O)C_{1-6}alkoxy$, N-morpholino and N-pyrrolidino; and

Ar is an aromatic or heteroaromatic group, unsubstituted or substituted with 1-4 substituents,

independently selected from OH, C₁₋₆alkyl, C₁₋₆alkoxy, NH₂, NH-C₁₋₆alkyl, N(C₁₋₆alkyl)(C₁₋₆alkyl), SH, S-C₁₋₆alkyl, NO₂, CF₃, OCF₃ and halo; and

p is 1-4.

39. (New) The compound according to claim 38, wherein R³ is selected from C₁₋₄alkylCO₂, C₁₋₄alkyl(C=O)NH, or C₁₋₄alkyl(C=O)N(C₁₋₄alkyl).

40. (New) The compound according to claim 39, wherein R³ is selected from CH₃CO₂, CH₃CONH, or CH₃CONCH₃.

41. (New) The compound according to claim 38, wherein R⁴ is selected from C(X)R⁵ and C(NH₂)=C(CN)₂.

42. (New) The compound according to claim 41, wherein R⁴ is C(X)R⁵.

43. (New) The compound according to claim 42, wherein X is selected from O and S.

44. (New) The compound according to claim 42, wherein R⁵ is selected from NH₂, OH, NH(CH₂)_pAr, NH(CH₂)_pOH and C₁₋₄alkoxy.

45. (New) The compound according to claim 44, wherein p is 1-3.

46. (New) The compound according to claim 44, wherein R⁵ is selected from NH₂, OH, NH(CH₂)_pAr, NH(CH₂)_pOH and OCH₃.

47. (New) The compound according to claim 46, wherein p is 1-2.

48. (New) The compound according to claim 38, wherein Ar is an unsubstituted phenyl group or a phenyl group substituted with 1-4 substituents optionally selected from OH, C₁₋₆alkyl, C₁₋₆alkoxy, NH₂, NH-C₁₋₆alkyl, N(C₁₋₆alkyl)(C₁₋₆alkyl), SH, S-C₁₋₆alkyl, NO₂, CF₃, OCF₃ and halo.

49. (New) The compound according to claim 46, wherein Ar is an unsubstituted phenyl group or a phenyl group substituted with 1-4 substituents independently selected from OH, C₁₋₆alkyl, C₁₋₆alkoxy, NH₂, NH-C₁₋₆alkyl, N(C₁₋₆alkyl)(C₁₋₆alkyl), SH, S-C₁₋₆alkyl, NO₂, CF₃, OCF₃ and halo.

50. (New) The compound according to claim 48, wherein Ar is an unsubstituted phenyl group or phenyl group substituted with 1-2 substituents independently selected from OH, C₁₋₄alkyl, C₁₋₄alkoxy, NH₂, NH-C₁₋₄alkyl, N(C₁₋₄alkyl)(C₁₋₄alkyl), SH, S-C₁₋₄alkyl, NO₂, CF₃, OCF₃ and halo.

51. (New) The compound according to claim 50, wherein Ar is an unsubstituted phenyl group or phenyl group substituted with 1-2 substituents independently selected from OH, OCH₃, NH₂, NHCH₃, N(CH₃)₂, SH, SCH₃, CF₃, OCF₃ and halo.

52. (New - Withdrawn) A method according to claim 33 wherein said cancer is a hematopoietic cell cancer.

53. (New - Withdrawn) A method according to claim 33 wherein said cancer is a leukemia, a lymphoma, a myeloma or a carcinoma.